

Message Text

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61

ORIGIN HEW-06

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E.O. 11652: N/A

TAGS: OGEN, TBIO, XX

SUBJECT: FDA ADVISORY - DEFECTIVE DEVICE

1. FDA ADVISES THAT THE COLEMAN 124, AUTOMATIC, 4 POSITION CELL
PROGRAMMER SPECTROPHOTOMETER, MANUFACTURED BY:

NAKA WORKS OF HITACHI LIMITED
882 ICHIGO, KATSUTA
IBARAKI, JAPAN

AND DISTRIBUTED BY:

PERKIN-ELMER, COLEMAN DIVISION
2000 YORK ROAD
OAK BROOK, ILLINOIS 60521

SINCE 1969 HAS AN ENGINEERING DEFECT IN THE FOUR POSITION TURRET ACCESSORY. THE ESSENTIAL PROBLEM IS THAT THE SERVICE AND OPERATION MANUALS PROVIDE INCOMPLETE DIRECTIONS FOR USE WHICH MAY RESULT IN IMPROPER INSTALLATION OF THE FOUR POSITION TURRET ACCESSORY ON THE INSTRUMENT. THIS MAY LEAD TO MISALIGNMENT OF THE LIGHT BEAM RESULTING IN ERRONEOUS DIAGNOSTIC READINGS AND UNRELIABLE SAMPLE RESULTS.

2. FOREIGN CONSIGNEES, AS FOLLOWS:

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